



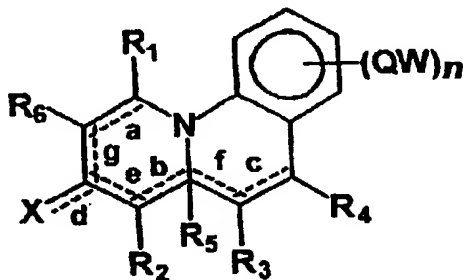
## INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification <sup>6</sup> : <b>A01N 43/90, C07D 455/02</b>		<b>A1</b>	(11) International Publication Number: <b>WO 99/05913</b>
			(43) International Publication Date: 11 February 1999 (11.02.99)
(21) International Application Number: PCT/EP98/04737 (22) International Filing Date: 29 July 1998 (29.07.98) (30) Priority Data: FI97A000193      1 August 1997 (01.08.97)      IT (71) Applicant (for all designated States except US): APPLIED RESEARCH SYSTEMS ARS HOLDING N.V. [NL/NL]; John B. Gorsiraweg 14, Curacao (AN). (72) Inventors; and (75) Inventors/Applicants (for US only): GUARNA, Antonio [IT/IT]; Via Pistoiese, 158, I-50040 Seano (Carmignano) (IT). SERIO, Mario [IT/IT]; Via di Baroncelli, 29, I-50012 Bagno a Ripoli (IT). (74) Agent: GERVASI, Gemma; Notarbartolo & Gervasi, Corso di Porta Vittoria, 9, I-20122 Milan (IT).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, GM, HR, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).  Published With international search report. Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.	

(54) Title: USE OF BENZO[C]QUINOLIZINE DERIVATIVES AS PLANT GROWTH REGULATORS

## (57) Abstract

Described herein is the use of benzo[c]quinolizine derivatives of formula (I) as regulators of the growth of plants, and compositions for agricultural use containing the said derivatives or their salts.



(I)

**FOR THE PURPOSES OF INFORMATION ONLY**

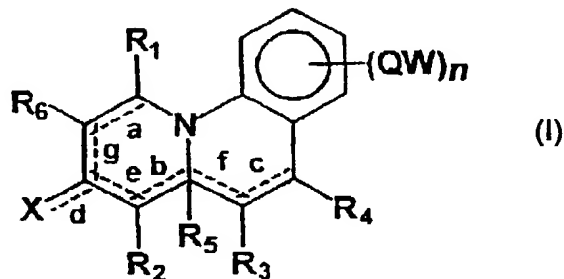
Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

## USE OF BENZO[C]QUINOLIZINE DERIVATIVES AS PLANT GROWTH REGULATORS

**Scope of invention**

- 5 The present invention regards the use of benzo[c]quinolizine derivatives of general formula (I)



in which:

- 15  $R_1$ ,  $R_2$ ,  $R_3$ ,  $R_4$ , and  $R_6$ , which are the same or different from one another, are chosen in the group consisting of: H,  $C_{1-8}$  alkyl,  $C_{2-8}$  alkenyl,  $C_{2-8}$  alkynyl,  $C_{3-6}$  cyclo-alkyl, aryl, heterocycle, halogen, CN, hazide, NRR',  $C_{1-8}$  alkylamine, arylamine,  $C_{1-8}$  alkyloxy, aryloxy, COOR, and CONRR', where R and R', which are the same or different from one another, are chosen in the group consisting of H,  $C_{1-8}$  alkyl, cyclo-alkyl, aryl, heterocycle, and aryl- $C_{1-8}$  alkyl;
- 20  $R_5$  is chosen in the group consisting of H,  $C_{1-8}$  alkyl, aryl- $C_{1-8}$  alkyl, COOR, CN, aryl, heterocycle, and the  $C_{1-8}$  alkyl heterocycle;
- X is chosen in the group consisting of O,  $C(=O)R$ , COOR,  $NO_2$ , and CONNR', in which R and R' are as defined above;
- 25 Q is chosen in the group consisting of: single-bond,  $C_{1-8}$  alkyl,  $C_{2-8}$  alkenyl,  $C_{2-8}$  alkynyl, cyclo-alkyl, CO, CONR, and NR, where R is as defined previously;
- W is chosen in the group consisting of H,  $C_{1-8}$  alkyl,  $C_{2-8}$  alkenyl,  $C_{2-8}$  alkynyl, cyclo-alkyl, trifluoromethyl,  $C_{1-8}$  alkoxy,  $C_{1-8}$  alkoxy- $C_{1-8}$  alkyl, aryl- $C_{1-8}$  alkyl, aryl, aryloxy, arylamine,  $C_{1-8}$  alkyl-carbonyl, arylcarbonyl, halogen, CN, NRR',  $C_{1-8}$  alkylamine, and heterocycle, in which the alkyl, alkenyl, alkynyl, cyclo-alkyl, aryl, and
- 30 heterocycle groups may be substituted;
- $n$  is 1, 2, 3 or 4;

the mark      indicates that the respective bonds a, b, c, d, e, f, and g may be single or double bonds, considering that when b or f are a double bond, the R<sub>5</sub> group is absent,

as regulators of the growth of plants.

#### 5   **State of the art**

It is known that steroidal enzymes have a considerable importance both in the field of medicine and in that of sciences related to the development of agriculture and foodstuffs. However, whilst the physiological role of steroids in man has been amply studied and documented, the physiological role of steroids in the vegetable  
10   world is less well known.

The growth of plants is governed by complex interactions between environmental signals and internal factors. Light regulates many processes of development throughout the life cycles of the plant, starting from seed germination right up to flower development (J. Chory, Trends, Genet., 1993, 9, 167). In the presence of  
15   light, the growth of hypocotyledons is inhibited, cotyledons expand, leaves develop, chloroplasts differentiate producing chlorophyll, and a large number of light-inducible genes are activated.

It has been recently suggested that brassinosteroids, which are the most widespread steroids in higher plants, may be directly involved in the response of  
20   plants to light (Chory *et al.*, Proc. Natl. Acad. Sci. 1996, 93, 12066). However, the interactions between phototransduction and hormones are not yet well known. Brassinolid is the steroid that has a fundamental role in the development of plants under the effect of light and has been identified in the larger part of higher plants. It is generated through a metabolic cascade in which campesterol (a compound  
25   similar to cholesterol) is reduced to campestanol. The enzyme which is responsible for this reduction is a 5-alpha-reductase steroidal enzyme, named DET2 after the gene of the same name isolated for the plant *Arabidopsis*, which presents a sequence analogy of up to 80% on the conservative amino acids of the iso-enzymes 1 and 2 of the 5-alpha-reductase of humans and rats.

30   The genetic mutations that inactivate DET2 do not allow production of brassinolid and determine deep alterations in the development of seeds and in *Arabidopsis* plants in the dark or under the effect of light.

In the dark, the plants mutated genetically in DET2 present short and thick hypocotyledons, accumulate anthocyanins, have open and expanded cotyledons, and develop primary-leaf buds. In the light, the mutated plants are smaller and of a darker green, have a reduced apical dominance and male fertility. In addition, they have different responses to light, with delayed flowering and delayed ageing of leaves and chlorophyll (J. Li *et al.*, Science, 1996, 272, 398).

These alterations of the mutating species are reversed with the exogenous addition of brassinolid to the growth medium (J. Li *et al.*, Proc. Natl. Acad. Sci., 1997, 94, 3554-3559).

#### 10 Detailed description of the invention

It has now been surprisingly found that the products of formula (I) as described above exert an inhibiting action on the 5-alpha-reductase steroidal enzymes, in particular on the DET2 enzyme, and hence are able to affect selectively the growth of plants in the dark and in the light, and can therefore be used as phytopharmaceutical substances in the field of agriculture and foodstuffs both as substances capable of improving the morphogenesis and development of plants that are commercially useful and as potential herbicides that inhibit the development of weeds.

In the products of formula (I) according to the present invention, by C<sub>1-8</sub> alkyl, C<sub>1-8</sub> alkenyl and C<sub>1-8</sub> alkynyl group are meant alkyl radicals, either linear or branched, such as methyl, ethyl, propyl, isopropyl, butyl, pentyl, hexyl, heptyl, octyl, ethylene, propene, butene, isobutene, acetylene, propyne, butyne, etc.

By the term cyclo-alkyl the following are meant: cyclopropane, cyclobutane, cyclopentane, cyclohexane, cycloheptane, cyclo-octane, norbornane, camphane, and adamantane.

By the term aryl the following are meant: phenyl and naphthyl.

By the term heterocycle the following are meant in particular: saturated or aromatic heterocycles containing one or more nitrogen atoms, and more in particular, pyridine, imidazole, pyrrole, indole, triazoles, pyrrolidine, and piperidine.

By halogen the following are meant: fluorine, chlorine, bromine, and iodine.

The substituents of the aforementioned W groups are preferably: halogen, OR, phenyl, NRR', CN, COOR, CONRR', and C<sub>1-8</sub> alkyl (in which R and R' are as defined above).

In particular, according to the present invention the products of formula (I) are preferred in which:

R<sub>5</sub> = H, heterocycle, aryl-C<sub>1-8</sub> alkyl, or C<sub>1-8</sub> alkyl heterocycle;

X = O;

Q = single-bond, CO, CONR, or NR (where R is as defined above);

W = H, F, Cl, Br, Me, *tert*-butyl, C<sub>1-8</sub> alkoxy, 2,5-dimethylhexyl, trifluoromethyl, 2,5-(di-trifluoromethyl)-phenyl, 4-methoxy-phenyl, 4-fluoro-phenyl, phenyl, phenyl-C<sub>1-8</sub> alkyl, C<sub>1-8</sub> alkylcarbonyl, or phenylcarbonyl;

n = 1 or 2;

R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub>, R<sub>6</sub> = H, Me, CN, phenyl, COOR, or CONRR' (where R and R' are as defined above).

Products preferred according to the present invention are:

1,2,4,4a,5,6 hexahydro-(11H)-benzo[c]quinolizin-3-one;

8-chloro-1,2,4,4a,5,6 hexahydro-(11H)-benzo[c]quinolizin-3-one;

1,2,4,4a,5,6 hexahydro-8-methyl(11H)-benzo[c]quinolizin-3-one;

1,2,4,4a,5,6 hexahydro-4-methyl-(11H)-benzo[c]quinolizin-3-one;

1,2,4,4a,5,6 hexahydro-1-methyl-(11H)-benzo[c]quinolizin-3-one;

1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

8-chloro-1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

8-methyl-1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

4-methyl-1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

1-methyl-1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

4a-benzyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

8-chloro-4a-benzyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

5,6-dihydro-(11H)benzo[c]quinolizin-3-one;

8-chloro-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

8-chloro-1-methyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

8-methyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

(*cis*) and (*trans*) 4-methyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;  
8-chloro-4-methyl-1,2,5,6-tetrahydro-(11H)-benzo[c]  
quinolizin-3-one;

4,8-dimethyl-1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

5 (*cis*) and (*trans*) 4,8-dimethyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

(*cis*) and (*trans*) 8-chloro-4-methyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one.

The products of formula (I) may be prepared according to known techniques. The products, and the processes for their preparation, are in any case described in the  
10 parallel Patent Application No. PCT/EP97/00552.

For this purpose, a number of products of formula (I) were tested as regards their capacity for modifying seed germination.

The introduction of compounds of formula (I) in various concentrations into the culture medium notably modified germination of *Arabidopsis Thaliana* Columbia  
15 (Col 0) seeds, kept in the dark for 10 days, as compared to the germination of non-treated seeds.

The effects observed on the treated seeds (i.e., lower growth of the hypocotyledons, development of cotyledons, and budding of the primary leaves) were similar to those described for seeds of plants genetically mutated as regards  
20 the DET2 enzyme, this indicating that the compounds of formula (I) are effective inhibitors in regard to this enzyme.

This observation indicates that the compounds of formula (I), in so far as they are inhibitors of the 5-alpha-reductases, and in particular of DET2 in plants, may be used to modify the germination of the seeds in the dark (if applied on the seeds)  
25 and modify the growth of the plants in the light (if applied on the plants). The possible industrial applications may thus regard the increase in germination of seeds of plants useful in agriculture and/or the reduction in the growth of harmful plants.

#### Example

30 Batches of 25 seeds of *Arabidopsis Thaliana* Columbia (Col 0) were made to germinate in a suitable culture medium consisting of 0.5 x MS at pH 5.7, containing 1% sucrose, 1x Vitamin B5 Gamborg, and 0.8% phyto-agar in the

presence of the 1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one inhibitor at the concentrations of 0, 0.01, 0.1, 1 and 10 micromolar. After two hours of treatment in light, the seeds were covered with three layers of aluminium foil and kept at 21°C in a growth chamber. After 10 days in the dark, the length of the  
5 hypocotyledons was measured. In the plants not treated with the inhibitor, the length measured was approximately 15-16 mm, whereas in the treated plants the hypocotyledons were progressively shorter as the concentration of inhibitor used was increased.

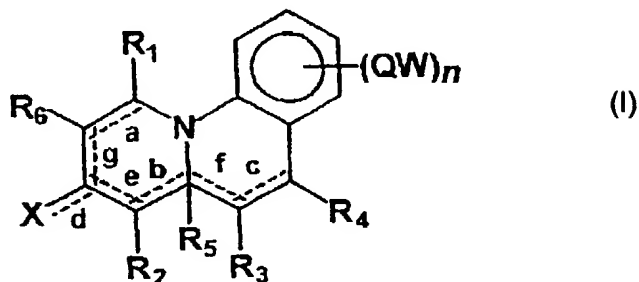
A reduction of 50% in the hypocotyledons as compared to the controls was found  
10 for the seeds treated with a concentration of 0.1 micromolar of inhibitor.

This indicates that the inhibitor at that concentration determines a control over germination of seeds in the dark.



## CLAIMS

1. Use of benzo[c]quinolizine compounds of general formula (I)



in which:

R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub>, R<sub>4</sub>, and R<sub>6</sub>, which are the same or different from one another, are chosen in the group consisting of H, C<sub>1-8</sub> alkyl, C<sub>2-8</sub> alkenyl, C<sub>2-8</sub> alkynyl, C<sub>3-8</sub> cyclo-alkyl, aryl, heterocycle, halogen, CN, hazide, NRR', C<sub>1-8</sub> alkylamine, arylamine, C<sub>1-8</sub> alkyloxy, aryloxy, COOR, and CONRR', where R and R', which are the same or different from one another, are chosen in the group consisting of H, C<sub>1-8</sub> alkyl, aryl, heterocycle, aryl-C<sub>1-8</sub> alkyl, and cyclo-alkyl;

R<sub>5</sub> is chosen in the group consisting of H, C<sub>1-8</sub> alkyl, aryl-C<sub>1-8</sub> alkyl, COOR, CN, aryl, heterocycle, and the C<sub>1-8</sub> alkyl heterocycle;

X is chosen in the group consisting of O, C(=O)R, COOR, NO<sub>2</sub>, and CONNR', in which R and R' are as defined above;

Q is chosen in the group consisting of single-bond, C<sub>1-8</sub> alkyl, C<sub>2-8</sub> alkenyl, C<sub>2-8</sub> alkynyl, cyclo-alkyl, CO, CONR, and NR, where R is as defined previously;

W is chosen in the group consisting of H, C<sub>1-8</sub> alkyl, C<sub>2-8</sub> alkenyl, C<sub>2-8</sub> alkynyl, cyclo-alkyl, trifluoromethyl, C<sub>1-8</sub> alkoxy, C<sub>1-8</sub> alkoxy-C<sub>1-8</sub> alkyl, aryl-C<sub>1-8</sub> alkyl, aryl, aryloxy, arylamine, C<sub>1-8</sub> alkyl-carbonyl, arylcarbonyl, halogen, CN, NRR', C<sub>1-8</sub> alkylamine, and heterocycle, in which the alkyl, alkenyl, alkynyl, cyclo-alkyl, aryl, and heterocycle groups may be substituted;

n is 1, 2, 3 or 4;

the mark      indicates that the respective bonds a, b, c, d, e, f, and g may be single or double bonds, considering that when b or f are a double bond, the R<sub>5</sub> group is absent; (or salts thereof),

as regulators of the growth of plants.

2. Use according to Claim 1, in which the inhibited 5-alpha-reductase steroidal enzyme is DET2.

3. Use according to Claims 1 and 2, in which in the products of formula (I):

5  $R_5 = H$  or heterocycle;

$X = 0$ ;

$Q = \text{single-bond, CO, CONR, or NR (where R is as defined above)}$ ;

$W = H, F, Cl, Br, Me, \text{tert-butyl, } C_{1-8} \text{ alkoxy, 2,5-dimethylhexyl, trifluoromethyl, 2,5-}$   
 (di-trifluoromethyl)-phenyl, 4-methoxy-phenyl, 4-fluoro-phenyl, phenyl, phenyl- $C_{1-8}$

10 alkyl,  $C_{1-8}$  alkylcarbonyl, or phenylcarbonyl;

$n = 1$  or  $2$ ;

$R_1, R_2, R_3, R_4, R_6 = H, Me, CN, \text{phenyl, COOR, or CONRR' (where R and R' are as defined above)}$ .

4. Use according to Claim 3, in which the products of formula (I) are:

15 1,2,4,4a,5,6 hexahydro-(11H)-benzo[c]quinolizin-3-one;

8-chloro-1,2,4,4a,5,6 hexahydro-(11H)-benzo[c]quinolizin-3-one;

1,2,4,4a,5,6 hexahydro-8-methyl(11H)-benzo[c]quinolizin-3-one;

1,2,4,4a,5,6 hexahydro-4-methyl-(11H)-benzo[c]quinolizin-3-one;

1,2,4,4a,5,6 hexahydro-1-methyl-(11H)-benzo[c]quinolizin-3-one;

20 1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

8-chloro-1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

8-methyl-1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

4-methyl-1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

1-methyl-1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

25 4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

5,6-dihydro-(11H)benzo[c]quinolizin-3-one;

8-chloro-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

8-chloro-1-methyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

8-methyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

30 (*cis*) and (*trans*) 4-methyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

8-chloro-4-methyl-1,2,5,6-tetrahydro-(11H)-benzo[c]  
 quinolizin-3-one;

4,8-dimethyl-1,2,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;  
(*cis*) and (*trans*) 4,8-dimethyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;  
(*cis*) and (*trans*) 8-chloro-4-methyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one.

- 5 5. Compounds of formula (I) as defined in Claim 1, in which:

$R_5 = C_{1-8}$ -alkyl-aryl or  $C_{1-8}$  alkyl heterocycle, and the other substituents are as defined in Claim 1.

6. Compounds according to Claim 5 of formula:

4a-benzyl-4,4a,5,6-tetrahydro-(11H)-benzo[c]quinolizin-3-one;

- 10 8-chloro-4a-benzyl-4,4a,5,6-tetrahydro-(11H)-benzo[c] quinolizin-3-one.

7. Composition for regulating the growth of plants, containing as active principle a product of formula (I) according to Claim 1 or mixtures of such products, possibly in combination with additives commonly used in agriculture for this type of preparations.

- 15 8. Process for regulating the growth of plants in which on the seeds and/or on the plants an effective quantity of a composition according to Claim 7 is distributed.

# INTERNATIONAL SEARCH REPORT

International Application No.

PCT/EP 98/04737

**A. CLASSIFICATION OF SUBJECT MATTER**  
IPC 6 A01N43/90 C07D455/02

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)  
IPC 6 A01N C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, Y	WO 97 29107 A (APPLIED RESEARCH SYSTEMS ; GUARNA ANTONIO (IT); SERIO MARIO (IT)) 14 August 1997 cited in the application see the whole document ---	1-8
Y	CHEMICAL ABSTRACTS, vol. 126, no. 25, 23 June 1997 Columbus, Ohio, US; abstract no. 327248, J. LI ET AL.: "Conservation of function between mammalian and plant steroid 5.alpha.-reductases" XP002086370 see abstract & PROC. NATL. ACAD. SCI. U.S.A., vol. 94, no. 8, 1997, pages 3554-3559, --- -/--	1-8



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

### \* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

1 December 1998

Date of mailing of the international search report

14/12/1998

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2  
NL - 2280 HV Rijswijk  
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,  
Fax: (+31-70) 340-3016

Authorized officer

Lamers, W

## INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 98/04737

## C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	CHEMICAL ABSTRACTS, vol. 124, no. 23, 3 June 1996 Columbus, Ohio, US; abstract no. 309107, J.LI ET AL.: "A role for brassinosteroids in light-dependent development of Arabidopsis" XP002086371 see abstract & SCIENCE, vol. 272, no. 5260, 1996, pages 398-401, ----	1-8
Y	DATABASE WPI Section Ch, Week 9519 Derwent Publications Ltd., London, GB; Class C03, AN 95-144764 XP002086372 & JP 07 069987 A (SANKYO CO LTD) , 14 March 1995 see abstract ----	1-8
X	R.M.ACHESON ET AL.: "Addition Reactions of Heterocyclic Compounds. Part XLV. New Azepines from Substituted 2-Methylquinolines and Dialkyl Acetylenedicarboxylates" JOURNAL OF THE CHEMICAL SOCIETY, SECTION C: ORGANIC CHEMISTRY., no. 19, 1971, pages 3291-3296, XP002086369 LETCHWORTH GB see page 3296, column 2, paragraph 2 -----	5

# INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/EP 98/04737

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 9729107 A	14-08-1997	IT FI960019 A	11-08-1997
		AU 1767297 A	28-08-1997
		CZ 9802289 A	14-10-1998
		NO 983444 A	24-07-1998
-----			